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## Valvular Heart Disease

### PROGNOSTIC VALUE OF MYOCARDIAL INJURY FOLLOWING UNCOMPLICATED TRANSCATHETER AORTIC VALVE IMPLANTATION

ACC Moderated Poster Contributions  
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Session Title: TAVR: Real World Outcomes and Potential Complications  
Abstract Category: 11. Valvular Heart Disease: Therapy  
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Authors: *Nazario Carrabba, Renato Valenti, Angela Migliorini, Guido Parodi, Alberto Santini, Ruben Vergara, David Antoniucci, Division of Cardiology, Careggi Hospital, Florence, Italy*

**Background:** Very few data exist about the clinical significance of myocardial injury (MI) occurring after transcatheter aortic valve implantation (TAVI) procedures. We sought to determine the incidence and the prognostic value of MI associated with uncomplicated TAVI.

**Methods:** Overall 68 consecutive patients (age  $81.1 \pm 6.1$  years, female 44%) who underwent TAVI procedure were included (Corevalve Medtronic; aortic valve area  $0.53 \pm 0.12$  cm<sup>2</sup>; n=65 transfemoral approach, n= 3 subclavian approach;). Patients who died within 24 hour following TAVI precluding cardiac biomarker measurements (n=3: 1 cardiac tamponade; 1 life-threatening arrhythmias; 1 TAVI failure) and those with major procedural complications were excluded (n=3: 1 ictus; 1 lung failure; 1 irreversible shock). Cardiac troponin I (cTnI) levels were determined at baseline, 6, 12, 24, 48, and 72 h following TAVI. Left ventricular ejection fraction (LVEF) were measured (Simpson method) at baseline, after TAVI, at 1, 6 and 12 months by 2D-echo. Acute kidney injury (AKI) was defined according Valve Academic Research Consortium (VARC).

**Results:** TAVI was associated with some degree of MI in 100% of the patients as determined by a rise in cTnI (median  $3.49 \pm 5.35$  ng/ml; range from 0.26 to 26.9 ng/ml). According VARC criteria, no peri-procedural myocardial infarction occurred. The development of AKI was found in 15% (9/62) of patients, and AKI was associated with a higher increase in biomarkers of MI ( $p = 0.008$ ). At 1 year follow-up: 4 patients died, and 8 patients were hospitalized for congestive heart failure. Overall a  $3.7 \pm 6.7\%$  improvement of LVEF was found following TAVI. The degree of MI (as peak cTnI) was not associated with a recovery of LVEF at 1 year follow-up ( $p = 0.981$ ). By Cox analysis, the development of AKI was an independent predictor of poor prognosis at 1-year (mortality and HF) (HR: 6.32, 95% CI 1.57 to 25.46,  $p < 0.009$ ), but not the degree of MI ( $p=0.578$ ).

**Conclusions:** In our center, TAVI (non transapical approach) was systematically associated with some degree of MI, with development of AKI determining a higher degree of MI. The development of AKI rather than the degree of MI was associated with a poor prognosis after uncomplicated TAVI.